

U.S.S.N. 10/623,398

Filed: July 18, 2003

AMENDMENT AND RESPONSE TO RESTRICTION REQUIREMENT**Amendment****In the Claims**

Please amend the claims as follows.

1. (previously presented) A method of using a glycosaminoglycan-degrading enzyme to reduce a cell proliferative response comprising

administering an effective amount of the enzyme to an individual in need thereof,

wherein the enzyme removes dermatan sulfate or chondroitin sulfate from cell surfaces thereby effectively decreasing growth factor receptors on the cell surfaces, and

wherein decreasing the growth factor receptors thereby decreases the cell proliferative response to such growth factors.
2. (Original) The method of claim 1 wherein the enzyme is selected from the group consisting of bacterial dermatan or chondroitin sulfate degrading enzyme and is selected from the group consisting of chondroitinase AC from *Flavobacterium heparinum*, chondroitinase B from *Flavobacterium heparinum*, chondroitin sulfate degrading enzymes from *Bacteroides* species, chondroitin sulfate degrading enzymes from *Proteus vulgaris*, chondroitin sulfate degrading enzymes from *Micrococcus*, chondroitin sulfate degrading enzymes from *Vibrio* species, chondroitin sulfate degrading enzymes from *Arthrobacter aureescens*, arylsulfatase B, N-acetylgalactosamine-6-sulfatase and iduronate sulfatase from mammalian cells, these enzymes expressed from recombinant nucleotide sequences in bacteria and combinations thereof.
3. (cancelled)
4. (Original) The method of claim 1 wherein the enzyme is a bacterial enzyme.

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5. (previously presented) The method of claim 4 wherein the enzyme is chondroitinase B.
6. (previously presented) The method of claim 1 wherein the enzyme is administered to decrease fibroblast proliferation.
7. (previously presented) The method of claim 1 wherein the chondroitin sulfate is selected from the group consisting of chondroitin sulfate B, chondroitin sulfate A and chondroitin sulfate C.
8. (previously presented) The method of claim 1 wherein the individual has scarring or is at risk of scarring.
9. (Original) The method of claim 1 wherein the enzyme is administered systemically.
10. (Original) The method of claim 1 wherein the enzyme is administered topically or locally at or adjacent to a site in need of treatment.
11. (Original) The method of claim 1 wherein the enzyme is administered in a controlled and/or sustained release formulation.
- 12-19. (cancelled)